

# **Study of drug treatments suitable for intravenous to oral switching**

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## BACKGROUND

Due to several factors intravenous drug treatments (IDT) in hospitals are not always changed to oral administration when possible.

## PURPOSE

To determine the frequency of IDT that can be switched to oral administration in a tertiary care hospital and to estimate savings due to switching the administration route of the selected drugs.

## **MATERIALS AND METHODS**

We collected prescription data on a randomly chosen weekday of all inpatients from hospital units with unit dose drug distribution. Four drugs with oral bioavailability greater than 75% were chosen for the study (Acetaminophen/Paracetamol, Levofloxacin, Omeprazole, Ranitidine).

Variables collected were: Prescribing service, medical specialty, type of diet, number of prescribed drugs administered orally, number of days of IDT and prescription of antiemetics.
Data was obtained from the pharmacy inpatient program (*Farmatools*<sup>®</sup>) and the hospital diet request system (*Dietools*<sup>®</sup>).

•A drug was considered for intravenous-to-oral switch therapy when the patient tolerated oral diet, had two or more drugs prescribed for oral administration, had received I.V. therapy for more than a day and had no antiemetic drugs prescribed.

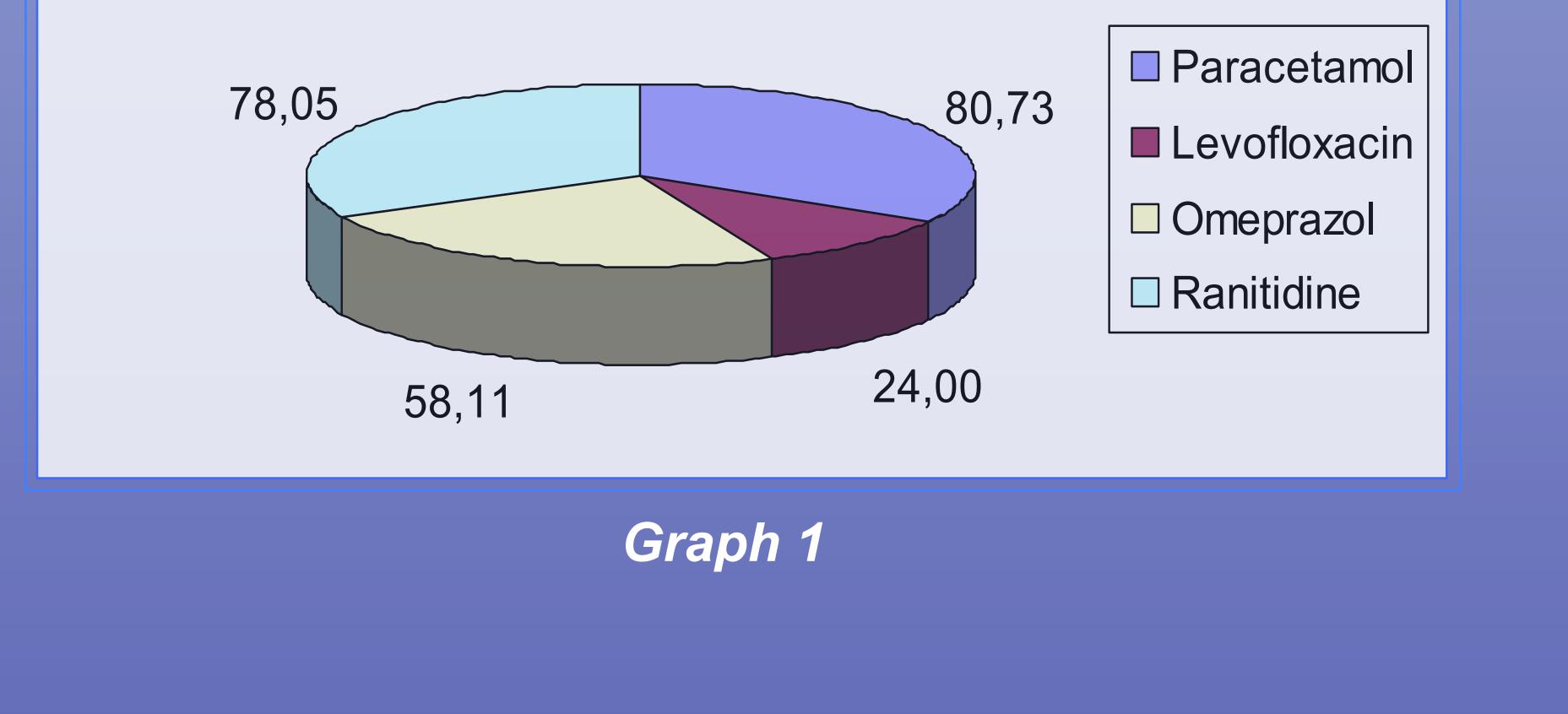
•Cost evaluation was based on drug prices obtained by the institution.

# •Prescriptions of 193 patients were analyzed: 169 of them were likely to be changed to oral administration. (% Prescriptions suitable for I.V. to oral switching: Graph 1).

•Estimated savings of Paracetamol prescriptions on the day of study added up to 296.45€, annual saving estimation of 108,204€. Savings due to levofloxacin would be 11.82€ (4,314€ per year), omeprazole 29.67€ (10,829€ per year) and ranitidine 12.93€ (4,719€ per year).

# % Prescriptions suitable for intravenous to oral switching

**CP-119** 



# CONCLUSIONS

Implementation and optimization of an intravenous-to-oral switch therapy program in a selected group of drugs would not only reduce complications associated with intravenous administration but also drug costs.

